

REQUEST FORM FOR
APPLICATION UNDER 37 CFR 1.53(b)

DOCKET NUMBER: 43890-427

Prior Application:

Art Unit: 2831

Examiner: E. Thomas

Assistant Commissioner for Patents
Washington, DC 20231

Sir:

This is a Request for filing a **Divisional** application under 37 CFR 1.53(b) of pending prior application

Serial No. 09/233,936, filed on January 20, 1999, entitled **ELECTROLYTIC CAPACITOR AND ITS**

MANUFACTURING METHOD, by the following named inventors: Kazuyo SAITO, Yukihiro NITTA, Hiroshi

TADA, Shigeyoshi IWAMOTO.

1. ☒ I hereby state that the enclosed application contains no new matter.
2. Oath or Declaration
 - a. ☐ Newly executed (original or copy)
 - b. ☒ Copy from a prior application (37 CFR 1.63(d))
 - i. ☐ Deletion of inventor(s)
Signed statement attached deleting inventor(s) named in the prior application, see 37 CFR 1.63(d)(2) and 1.33(b).
3. ☒ Incorporation By Reference (useable if Box 2b is checked)
The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 2b, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.
4. ☐ Preliminary Amendment is enclosed.
5. ☒ An Information Disclosure Statement and PTO1449 Form are submitted herewith.
6. ☒ Cancel claims 1-20.

3c490 U.S. PTO
09/616944
07/14/00

3c781 U.S. PTO
07/14/00

3c781 U.S. PTO
07/14/00

7. The filing fee is calculated on the basis of the claims existing in the prior application as amended at 4 and 6 above:

	NO. OF CLAIMS		EXTRA CLAIMS	RATE	AMOUNT
Total Claims	12	+20	0	\$18.00 =	\$0.00
Independent Claims	1	-3	0	\$78.00 =	\$0.00
Basic Application Fee					\$690.00
If multiple dependent claims are presented, add \$0.00					\$0.00
Total Application Fee					\$690.00
Subtract ½ if small entity					\$0.00
TOTAL APPLICATION FEE DUE					\$690.00
AMOUNT TO BE CHARGED TO DEPOSIT ACCOUNT NO. 500417					\$690.00

- 7a. ☐ Enclosed is a Verified Statement to establish small entity status under 37 CFR 1.9 and 37 CFR 1.27.
- 7b. ☐ A verified Statement to establish small entity status under 37 CFR 1.9 and 37 CFR 1.27 was filed in prior application and such status is still proper and desired.
- 8a. ☒ **PLEASE CHARGE DEPOSIT ACCOUNT 500417 in the amount of \$690.00**
- 8b. ☒ The Commissioner is hereby authorized to charge fees under 37 CFR 1.16 and 1.17 which may be required, including any extension of time fees to maintain the pendency of the parent application Serial No. 09/233,936 or credit any overpayment to Deposit Account No. 500417.
9. ☒ Amend the specification by inserting before the first line the sentence:
 --This application is a Divisional of Application Serial No. 09/233,936 filed January 20, 1999--
10. ☒ Priority of Application Serial No. 10-015269, filed on January 28, 1998 in Japan, and Application Serial No. 10-350072, filed on December 9, 1998 in Japan, is claimed under 35 USC 119. The certified priority documents were filed in Serial No. 09/233,936 on April 12, 1999.
11. ☒ The prior application is assigned of record to
MATSUSHITA ELECTRIC INDUSTRIAL CO., LTD.
 Osaka, Japan
12. ☒ The power of attorney in the prior application is to:
 McDermott, Will & Emery
13. ☐ Also enclosed:

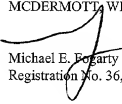
14. ☐ A petition, fee and response has been filed to extend the term in the pending prior application until .

Address all future communications to: (May only be completed by applicant, or attorney or agent of record)

McDermott, Will & Emery
600 13th Street, N.W.
Washington, DC 20005-3096

Respectfully submitted,

MCDERMOTT, WILL & EMERY


Michael E. Fogarty
Registration No. 36,139

600 13th Street, N.W.
Washington, DC 20005-3096
(202) 756-8000 MEF:dtb
Date: July 14, 2000
Facsimile: (202) 756-8087

ELECTROLYTIC CAPACITOR AND ITS MANUFACTURING METHOD

FIELD OF THE INVENTION

The present invention relates to an electrolytic capacitor as an electronic component, and its manufacturing method.

BACKGROUND OF THE INVENTION

In the recent trend of electronic appliances becoming digital and higher in frequency, the electrolytic capacitor, one of the electronic components, is required to be larger in capacity than in the conventional part and is superior in impedance characteristic in high frequency region. To meet such demand, it has been attempted to enhance the conductivity of the driving electrolyte (hereinafter called electrolyte), decrease the resistance of separator, or use a conductive compound obtained by making conductive a sheet insulator such as paper, cloth, nonwoven cloth or high polymer film, as a separator.

Also as an attempt to make the separator conductive, various methods have been proposed, such as kneading or mixing of carbon fibers or particles, and compounding with graphite powder. Moreover, by using monomer such as

pyrrole, thiophen or aniline, a method of forming a conductive high polymer on the surface by chemical oxidation and polymerization is disclosed (see Japanese Laid-open Patent No. 64-90517).

In the conventional constitution, however, there was a limit in enhancement of conductivity of electrolyte, and its conductivity is at most about ten to scores of mS/cm at the present, and electrolyte having a sufficient conductivity is not developed yet, and an electrolytic capacitor of an electrolyte having a sufficient impedance characteristic is not obtained so far.

On the other hand, for decrease of resistance of separator, it has been attempted to decrease the separator thickness, lower the density, make uniform the pore size, or change from paper to high polymer nonwoven fabric, but due to lack of strength by lowering of density and other problems, a sufficient effect of lowering the resistance is not obtained yet.

Further, the separator made conductive by kneading or mixing carbon fibers or particles is not sufficient in the electric conductivity, and it was hard to obtain a separator of low density. When using a separator by compounding graphite powder, there was a problem of increase of shorting due to drop of graphite powder and dispersion into electrolyte.

On the other hand, in the method of forming a conductive high polymer on the surface by chemical oxidation and polymerization from monomer of pyrrole, thiophen or aniline, it is difficult to compose an electrolytic capacitor of which rated voltage exceeds 35 V because there is almost no effect of deterioration of dielectric oxide film by oxidizing agent or chemical formation of conductive high polymer (defect repairing capability of dielectric oxide film). If composed, however, increase of leak current or shorting between anode and cathode may occur during agent process or high temperature test.

It is hence an object of the invention to present an electrolytic capacitor of high dielectric strength excellent in impedance characteristic, leak current property and reliability.

SUMMARY OF THE INVENTION

An electrolytic capacitor of the invention comprises:

(a) a capacitor element having an anode, a cathode, and a solid organic conductive material disposed between the anode and the cathode,

(b) an electrolyte,

(c) a case for accommodating the capacitor element and the electrolyte, and

(d) a sealing member disposed to cover the opening of the case.

A manufacturing method of electrolytic capacitor of the invention comprises:

- (a) a step of fabricating an anode,
- (b) a step of fabricating a cathode,
- (c) a step of forming a solid organic conductive material on the surface of the anode, and
- (d) a step of disposing an electrode between the anode having the solid organic conductive material and the cathode.

In this constitution, an electrolytic capacitor having an excellent impedance characteristic, an excellent leak current characteristic, an excellent reliability, and a high dielectric strength is obtained.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 (a) is a partial sectional perspective view showing a constitution of an electrolytic capacitor according to a first embodiment of the invention.

Fig. 1 (b) is a schematic diagram magnifying the essential parts of the electrolytic capacitor element shown in Fig. 1 (a).

Fig. 2 is a manufacturing process diagram for manufacturing the anode foil of an electrolytic capacitor

of the invention.

REFERENCE NUMERALS

- 1 Anode foil
- 2 Solid organic conductive material
- 3 Cathode foil
- 5 Anode lead
- 6 Cathode lead
- 7 Sealing member
- 8 Metallic case
- 9 Outer tube
- 10 Electrolyte
- 11 Dielectric oxide film
- 21 Aluminum foil
- 22 Etching foil
- 23 Solution containing polymerizable monomer
- 24 Heating oven

DETAILED DESCRIPTION OF THE INVENTION

An electrolytic capacitor of the invention comprises a case having an opening, an electrolyte contained in the case, a capacitor element placed in the electrolyte, and a sealing member disposed to cover the opening. The capacitor element has an anode, a cathode, and a solid organic conductive material installed between the anode and

cathode.

In this constitution, by making use of the high electric conductivity of the solid organic conductive material, the interpolar resistance in the conductive portion can be extremely decreased, and hence the impedance characteristic is enhanced. Further, by using together with the electrolyte having the repair capability of dielectric oxide film of valve action metal, an electrolytic capacitor of low leak current having a high dielectric strength is obtained.

In the invention, as the solid organic conductive material, an organic semiconductor is preferred, and preferred examples of organic semiconductor include 7,7,8,8-tetra-cyanoquinodimethane complex and its derivatives (hereinafter called TCNQ complexes). In this constitution, by dissolving and impregnating TCNQ complexes, a layer of solid organic conductive material having a high conductivity may be filled sufficiently to the inside of the pit of the anode processed by etching. As a result, an electrolytic capacitor excellent in impedance characteristic particularly in a high frequency region over 100 kHz can be obtained. Still more, TCNQ complexes can be directly applied on a separator base material. Moreover, a capacitor element winding an anode foil and a cathode foil through a separator base material can be manufactured.

Alternatively, a capacitor element laminating one set or two sets or more of anode and cathode through separator base material may be impregnated in a heated and dissolved TCNQ complex solution, and by cooling and solidifying, the conductivity may be easily expressed. As a result, an electrolytic capacitor having an excellent impedance characteristic may be easily obtained.

In the invention, the word of "anode" means "positive electrode", and the word of "cathode" means "negative electrode". The word of "conductive high polymer" means "conductive polymer".

In the invention, the conductive high polymer includes pyrrole, aniline, thiophen, ethylene dioxythiophen, sulfonated aniline, sulfonated pyrrole, sulfonated thiophen, sulfonated ethylene dioxythiophen, their derivatives, and polymers of various polymerizable monomers.

Methods of forming such polymers include a method by liquid-phase chemical polymerization, a method by vapor-phase chemical polymerization, a method by liquid-phase electrolytic polymerization, and a method by drying soluble high polymer solution and utilizing residual high polymer.

Usable examples of conductive high polymer include polypyrrole, polyethylene dioxythiophen, or polyaniline formed by chemical polymerization or electrolytic polymerization, and dry residual sulfonated polyaniline

obtained by drying solutions of soluble polyanilines.

In liquid-phase polymerization, in a solution containing at least the polymerizable monomer and a proper oxidizing agent, a capacitor element is immersed, and polymerized. In the case of an electrolytic polymerization, in a solution containing at least the polymerizable monomer and a proper oxidizing agent, a capacitor element is immersed, and power is supplied to polymerize it. In the case of vapor-phase polymerization, in a solution containing at least a proper oxidizing agent, a capacitor element is immersed (or immersed, lifted and dried), and then in the vapor phase containing at least the polymerizable monomer, the capacitor element is placed. By these methods, a layer of solid organic conductive material having a high conductivity can be sufficiently applied into the inside of pits of the anode processed by etching. As a result, an electrolytic capacitor having an excellent impedance characteristic also in a high frequency region of over 100 kHz in particular is obtained. Moreover, by polymerizing the conductive high polymer directly in the separator base material, anode or cathode in vapor phase, the conductivity can be easily expressed. As a result, an electrolytic capacitor having an excellent impedance characteristic may be easily obtained.

Further, since these conductive high polymers have a

high compatibility in an electrolyte composed of an organic matter, when impregnated with the electrolyte, it is quickly swollen and diffused into the inner parts of the conductive high polymer. Accordingly, when composing a capacitor element having the dielectric oxide film coated with conductive high polymer, the capacity for restoring the dielectric oxide film can be maintained at high level.

The electrolyte comprises an electrolytic substance such as salt of organic acid or salt of inorganic acid, and a solvent for dissolving such electrolytic substances. As such organic solvent, an organic solvent capable of swelling an organic conductive material by immersing it is preferred. In this constitution, as mentioned above, the electrolyte quickly swells and diffuses into the inner part of the conductive high polymer. Therefore, when composing a capacitor element having the dielectric oxide film coated with conductive high polymer, the capacity for restoring the dielectric oxide film can be maintained at high level.

As the electrolytic substance to be dissolved in the electrolyte, in a base for composing the electrolyte, when the concentration of the base or hydroxide of base is 1 wt.% and the measuring temperature is 30°C, the hydrogen ion concentration in aqueous solution of base or hydroxide of base is usable at 1.0×10^{-13} mol/dm³ or more. As its specific example, as the base for composing the

electrolytic substance, at least one is used as being selected from the group consisting of compound having alkyl substituent amidine group, quaternary salt of compound having alkyl substituent amidine group, tertiary amine and ammonium. In this constitution, leak of electrolyte is prevented, and a capacitor of enhanced reliability is obtained.

When composing a capacitor by using an electrolyte composed of an electrolytic substance of a base of a strong basicity of which hydrogen ion concentration is less than 1.0×10^{-13} mol/dm³ (for example, tetra-alkyl ammonium or tetra-alkyl phosphonium), in long-term environmental test in the compound environments of high temperature and high humidity (for example, 60°C and 95% RH), the sealing member is likely to be damaged by the effects of the base of strong basicity, and leak is likely to occur, and the reliability is slightly inferior.

As the base for composing the electrolytic substance, at least one is preferred to be used as being selected from the group consisting of compound having alkyl substituent amidine group, quaternary salt of compound having alkyl substituent amidine group, tertiary amine and ammonium. In these electrolytic substances, when the concentration of the base or hydroxide of base is 1 wt.% and the measuring temperature is 30 °C, the hydrogen ion concentration in

aqueous solution of base or hydroxide of base is 1.0×10^{-13} mol/dm³ or more. Accordingly, the leak due to such strong acidity as mentioned above is less likely to occur.

As the quaternary salt of compound having alkyl substituent amidine group, a quaternary compound formed by alkyl group or aryl alkyl group with 1 to 11 carbon atoms is preferred, and such compound is one selected from the group consisting of imidazole compound, benzoimidazole compound, and alicyclic amidine compound. In this constitution, when hydroxide ions are formed by electrolytic reaction in the electrolyte, since the reaction between hydroxide ion and amidine group of N-C-N or reaction of decomposition and ring opening is fast, the electrolytic products disappear quickly. As a result, even in the condition of high temperature and high humidity, leak of electrolyte to outside can be prevented.

The quaternary salt of compound having alkyl substituent amidine group is at least one selected from the group consisting of 1-methyl-1,8-diazabicyclo [5,4,0] undecene-7, 1-methyl-1,5-diazabicyclo [4;3,0] nonene-5, 1,2,3-trimethyl imidazolinium, 1,2,3,4-tetramethyl imidazolinium, 1,2-dimethyl-3-ethyl-imidazolinium, 1,3,4-trimethyl-2-ethyl imidazolinium, 1,3-dimethyl-2-heptyl imidazolinium, 1,3-dimethyl-2-(-3' heptyl) imidazolinium, 1,3-dimethyl-2-dodecyl imidazolinium, 1,2,3-trimethyl-

1,4,5,6-tetrahydropyrimidium, 1,3-dimethyl imidazolium, 1-methyl-3-ethyl-imidazolium, and 1,3-dimethyl benzoimidazolium. According to this constitution, the conductivity of the electrolyte can be heightened and an excellent heat resistance is realized. Therefore, external leak of electrolyte at high temperature and high humidity is prevented, and an electrolytic capacitor having an excellent high temperature stability and low impedance is obtained.

The boiling point of the solvent of the electrolyte is 200°C or more, the conductivity at measuring temperature of 30°C of the electrolyte is 1.0 mS/cm or more, and the spark ignition voltage is 80 V or more. In this constitution, it is effective to prevent the problem of electrolytic capacitor for surface mounting, that is, deformation of appearance due to elevation of capacitor internal pressure caused by heat treatment during surface mounting (both capacitor and substrate exposed to high temperature of soldering). Moreover, since the boiling point of the solvent of the electrolyte is high (vapor pressure is low), defective soldering when mounting is less likely to occur, and further since the conductivity is high, the impedance performance is maintained. Still more, since the spark ignition voltage is sufficiently high, an electrolytic capacitor having a high dielectric strength is obtained.

The sealing member is composed by using an elastic rubber. In the solid electrolytic capacitor for surface mounting using this elastic rubber, the adsorbed moisture contained in the electrode foil of capacitor element, separator, solid electrolytic substance and rubber sealing member, or the adsorbed moisture bonded to the inside of the case is evaporated at once at high temperature at the time of surface mounting. As a result, the pressure elevation inside the capacitor is extreme, which may lead to defective air tightness of the capacitor or scattering of sealing member. However, by containing the electrolyte using a solvent of high boiling point (low vapor pressure), the total pressure inside the capacitor can be lowered at the time of mounting. Therefore, by containing the liquid component (herein an electrolytic solution) in addition to the electrolytic substance of solid electrolytic substance type, pressure elevation inside the capacitor can be suppressed, and defective soldering and others can be improved.

Examples of solvent of electrolyte having a boiling point of 200°C or more include 3-alkyl-1,3-oxazolidine-2-one (more specifically, 3-methyl-1,3-oxazolidine-2-one: boiling point 260°C), 1,3-dialkyl-2-imidazolidinone other than 1,3-dimethyl-2-imidazolidinone (more specifically, 1,3-dimethyl-2-imidazolidinone: boiling point 236°C, 1,3-

diisopropyl-2-imidazolidinone: boiling point 255°C, 1-methyl-3-ethyl-2-imidazolidinone: boiling point 230 °C), 1,3,4-trialkyl-2-imidazolidinone (more specifically, 1,3,4-trimethyl-2-imidazolidinone: boiling point 241°C), 1,3,4,5-tetra-alkyl-2-imidazolidinone (more specifically, 1,3,4,5-tetramethyl-2-imidazolidinone: boiling point 249°C), cyclic lactone (more specifically, γ -butyrolactone: boiling point 204 °C), polyhydric alcohol (more specifically, ethylene glycol: boiling point 201°C, glycerin: boiling point 290°C), carbonate (more specifically, ethylene carbonate: boiling point 238°C, propylene carbonate: 242°C), and others.

Hereinafter, preferred embodiments of the invention and prior arts as comparative examples are described below while referring to the attached drawings.

Fig. 1 (a) and Fig. 1 (b) are a partial sectional perspective view showing a constitution of an electrolytic capacitor of the invention, and a conceptual view magnifying essential parts of its element. In Fig. 1 (b), the surface is roughened by etching process, and then a dielectric oxide film 11 is formed by oxidation treatment. An anode foil 1 composed of an aluminum foil forming a solid organic conductive material 2 on its surface, and a cathode foil 3 formed by etching an aluminum foil are wound around a separator 4. Or, after roughening the surface by etching process, an anode foil 1 composed of an aluminum

foil forming a dielectric oxide film 11 by oxidation treatment and a cathode foil 3 formed by etching process of aluminum foil are wound through an electrolytic paper 4A. Then by high temperature treatment thereof, the electrolytic paper 4A is treated by any method of carbonization treatment, and a capacitor element 12 or 12A is formed. Between the dielectric oxide film 11 and cathode foil 3, a solid organic conductive material 2 is formed. It is impregnated with an electrolyte 10, and swollen and infiltrated into the solid organic conductive material 2. Thus, the capacitor element 12 or 12A is composed. The capacitor element 12 or 12A is put in a cylindrical aluminum metallic case 8 with a bottom as shown in Fig. 1 (a). Further, the releasing end of the aluminum metallic case 8 is sealed so that a sealing member 7 made of rubber may penetrate through an anode lead 5 and a cathode lead 6 for external lead-out being led out from the anode foil 1 and cathode foil 3 from the sealing member 7. Thus, the side surface of the metallic case 8 is covered with an external tube 9.

Fig. 2 (a) to Fig. 2 (g) show the manufacturing process for manufacturing the anode foil 1 for electrolytic capacitor of the invention in batch. As shown in Fig. 2 (a), an etching foil 22 (Fig. 2 (b)) obtained by etching an aluminum foil 21 is oxidized. In this way, an anode foil 1

forming a dielectric oxide film 11 is formed (Fig. 2 (c)). Successively, this anode foil 1 is impregnated in a solution 23 containing a polymerizable monomer capable of forming a conductive high polymer layer as shown in Fig. 2 (d), and lifted, then heated (also dried) by a heating oven 24 as shown in Fig. 2 (e). Thus, as shown in Fig. 2 (f), an anode foil 1 forming a solid organic conductive material 2 on the surface is composed. Next, as shown in Fig. 1 (g), thus constituted anode foil 1 and the cathode foil 3 formed by etching the aluminum foil 21 are wound through a separator 4. In this way, a capacitor element 12 is composed. The subsequent process is same as in the above manufacturing method. The capacitor element 12 is put into a cylindrical metallic case 8 with a bottom together with electrolyte 10. The releasing end of the metallic case 8 is sealed, by using a sealing member 7, so that an anode lead 5 and a cathode lead 6 for external lead-out being led out from the anode foil 1 and cathode foil 3 respectively may penetrate through the sealing member 7. Thus, the side of the metallic case 8 is covered with an external tube 9.

The electrolyte used in the electrolytic capacitor of the invention is specifically described below.

As the solvent for the electrolyte of the electrolytic capacitor of the invention, an organic solvent stable electrically and chemically and capable of swelling in the

organic conductive material is used. Such organic solvent is desired to have a boiling point of 200°C or more. The solvent is preferred to be mainly composed of γ -butyrolactone and/or ethylene glycol. In addition, for the purpose of improving the low temperature characteristic and enhancing the dielectric strength, other organic solvent compatible with γ -butyrolactone and/or ethylene glycol may be mixed as a subsidiary solvent. Nevertheless, the subsidiary solvent is not required to be an organic solvent capable of swelling in an organic conductive material.

As the subsidiary solvent, in addition to the organic solvent with boiling point of 200°C or more mentioned above, the following organic solvents may be used either alone or as a mixed solvent of two or more kinds, that is, polyhydric alcohol system solvents (propylene glycol, diethylene glycol, 1,4-butane diol, polyoxy alkylene polyol), lactone system solvents (γ -valerolactone, δ -valerolactone, 3-ethyl-1,3-oxazolidine-2-one), water, amide system solvents (N-methyl formamide, N,N-dimethyl formamide, N-methyl acetamide), ether system solvents (methylal, 1,2-dimethoxy ethane, 1-ethoxy-2-methoxy ethane, 1,2-diethoxy ethane), nitrile system solvents (acetonitrile, 3-methoxy propionitrile), furane system solvents (2,5-dimethoxy tetrahydrofuran), 2-imidazolidinone system solvents (1,3-dimethyl-2-imidazolidinone), and others.

In the case of mixed solvent, the mixing ratio of the solvent is preferred to be 40 parts by weight of a solvent of which boiling point is less than 200°C in 100 parts by weight of solvent of which boiling point is 200°C or more. If the content of the solvent of which boiling point is less than 200°C is more than 40 parts, the heat resistance is lowered when an electrolytic capacitor is formed for surface mounting, and the defective rate of soldering becomes higher.

Examples of tertiary amine used in the electrolyte of the invention include trialkylamines (trimethylamine, dimethylethylamine, methyldiethylamine, triethylamine, dimethyl n-propylamine, dimethyl isopropylamine, methyl ethyl n-propylamine, methylethyl isopropylamine, diethyl n-propylamine, diethyl isopropylamine, tri-n-propylamine, tri-isopropylamine, tri-n-butylamine, tri-tert-butylamine, etc.), and phenyl group containing amines (dimethyl phenylamine, methylethyl phenylamine, diethyl phenylamine, etc.).

Among them, trialkylamine having a high conductivity is preferred. More preferably, it is preferred to use at least one selected from the group consisting of trimethylamine, dimethylethylamine, methyl diethylamine, and triethylamine, and by the use thereof, a capacitor high in conductivity and having an excellent impedance

performance is obtained.

Examples of compound having alkyl substituent amidine group used in the electrolyte of the invention include imidazole compound, benzoimidazole compound, and alicyclic amidine compound (pyrimidine compound, imidazoline compound). More specifically, it is preferred to use 1,8-diazabicyclo [5,4,0] undecene-7, 1,5-diazabicyclo [4,3,0] nonene-5, 1,2-dimethyl imidazolinium, 1,2,4-trimethyl imidazoline, 1-methyl-2-ethyl-imidazoline, 1,4-dimethyl-2-ethyl imidazoline, 1-methyl-2-heptyl imidazoline, 1-methyl-2-(3' heptyl) imidazoline, 1-methyl-2-dodecyl imidazoline, 1,2-dimethyl-1,4,5,6-tetrahydropyrimidine, 1-methyl imidazole, and 1-methyl benzoimidazole, and when these compounds are used, a capacitor having a high conductivity and excellent impedance performance is obtained.

In the examples of quaternary salt of compound having alkyl substituent amidine group used as the electrolyte is preferred to be a quaternary compound formed by alkyl group or aryl alkyl group with 1 to 11 carbon atoms is preferred, and preferred examples of amidine group are imidazole compound, benzoimidazole compound, and alicyclic amidine compounds (pyrimidine compound, imidazoline compound). Specific preferred examples include 1-methyl-1,8-diazabicyclo [5,4,0] undecene-7, 1-methyl-1,5-diazabicyclo [4,3,0] nonene-5, 1,2,3-trimethyl imidazolinium, 1,2,3,4-

tetramethyl imidazolinium, 1,2-dimethyl-3-ethyl-imidazolinium, 1,3,4-trimethyl-2-ethyl imidazolinium, 1,3-dimethyl-2-heptyl imidazolinium, 1,3-dimethyl-2-(-3' heptyl) imidazolinium, 1,3-dimethyl-2-dodecyl imidazolinium, 1,2,3-trimethyl-1,4,5,6-tetrahydropyrimidium, 1,3-dimethyl imidazolium, 1-methyl-3-ethyl-imidazolium, and 1,3-dimethyl benzoimidazolium. By using these compounds, it is possible to obtain an electrolytic capacitor with an excellent long-term stability having a high heat resistance, high conductivity, and excellent impedance performance.

Examples of organic acid used in the electrolyte of the invention include the following compounds: polycarboxylic acid (valence of 2 to 4), aliphatic polycarboxylic acid (saturated polycarboxylic acid, for example, oxalic acid, malonic acid, succinic acid, glutaric acid, adipic acid, pimelic acid, suberic acid, azelaic acid, sebacic acid, 1,6-decanoic dicarboxylic acid, 5,6-decanoic dicarboxylic acid, 1,7-octanoic dicarboxylic acid, and unsaturated polycarboxylic acid, for example, maleic acid, fumaric acid, itaconic acid, aromatic polycarboxylic acid (for example, phthalic acid, isophthalic acid, terephthalic acid, trimellitic acid, pyromellitic acid), alicyclic polycarboxylic acid (for example, cyclohexane-1,2-dicarboxylic acid, cyclohexene-1,2-dicarboxylic acid, etc.), hexahydrophthalic acid, alkyl substituents with 1 to 3

carbon atoms of their polycarboxylic acids (for example, citraconic acid, dimethyl maleic acid), or nitro substitutes (nitrophthalic acid, 3-nitrophthalic acid, 4-nitrophthalic acid), and polycarboxylic acid containing sulfur (for example, thiopropionic acid), monocarboxylic acid, aliphatic monocarboxylic acid with 1 to 30 carbon atoms (for example, formic acid, acetic acid, propionic acid, butyric acid, isobutyric acid, valeric acid, caproic acid, enanthic acid, caprylic acid, perargonic acid, lauric acid, mystyric acid, stearic acid, behenic acid, other saturated carboxylic acid, and acrylic acid, methacrylic acid, oleic acid, and other unsaturated carboxylic acid), aromatic monocarboxylic acid (for example, benzoic acid, o-nitrobenzoic acid, p-nitrobenzoic acid, cinnamic acid, naphthoic acid), and oxycarboxylic acid (for example, salicylic acid, mandelic acid, resorcylic acid). Among these compounds, particularly preferred compounds are those having high conductivity and excellent thermal conductivity such as maleic acid, phthalic acid, cyclohexane carboxylic acid, cyclohexene-1,2-dicarboxylic acid, cyclohexene-1,2-dicarboxylic acid, adipic acid, and benzoic acid.

The ratio of organic acid and base for composing the electrolyte is usually 4 to 11 at the pH of the electrolyte, and preferably 6 to 9. Out of this range, the spark voltage of the electrolyte (dielectric strength) is lowered.

As the electrolytic salt, organic carboxylic acids stable electrically and chemically are desired. Preferred examples of such organic carboxylic acid include maleic acid, phthalic acid, cyclohexane carboxylic acid, cyclohexene-1,2-dicarboxylic acid, cyclohexene-1,2-dicarboxylic acid, adipic acid, and quaternary salt of compound having alkyl substituent amidine group of benzoic acid.

The electrolyte of the electrolytic capacitor of the invention may mix and contain various additives as required. Usable additives include phosphor compounds (phosphoric acid, ester phosphate, etc.), boric acid compounds (boric acid, complex of boric acid and polysaccharides (mannite, sorbit, etc.)), nitro compounds (o-nitrobenzoic acid, n-nitrobenzoic acid, p-nitrobenzoic acid, o-nitrophenol, m-nitrophenol, p-nitrophenol, p-nitroacetophenone, etc.), and others. In the aluminum electrolytic capacitor, mixing of these additives will improve the restoration of aluminum oxide film. As a result, it is preferred because an electrolytic capacitor of high dielectric strength can be formed easily.

As the bar element for the terminal of the electrolytic capacitor of the invention, a material undergoing corrosion preventive treatment may be used. By corrosion preventive treatment of the bar element, electrolytic current can be

suppressed, and the sealing performance may be enhanced. Corrosion preventive treatment of the bar element is preferred to be done on both terminals of anode and cathode, but only either one may be treated. As means of corrosion preventive treatment, anode oxidation treatment in aqueous solution, coating-sintering of metal alkoxide, and coating-sintering of colloidal solution of metal oxide (colloidal solution of silicon dioxide and titanium dioxide) are convenient and preferred.

The sealing member 7 is preferably an elastic material mainly composed of rubber polymer formed of a copolymer of isobutylene, isoprene and divinyl benzene, comprising 0.5 to 20 parts of vulcanizing agent such as peroxide or alkyl phenol formalin resin. In other vulcanizing method using other vulcanizing agent than peroxide or alkyl phenol formalin resin (for example, sulfur vulcanization), the rubber elasticity drops significantly when left over for a long period in the condition of high temperature and high humidity, and sufficient sealing performance is not obtained, and as a result the organic conductive material may oxidize and deteriorate due to invasion of water from outside.

Preferred embodiments of the invention are described below. In the embodiments, a "part" always refers to a "part by weight."

The composition of the electrolytes used in the embodiments of the invention and in comparative examples is as follows. As the index of hydrogen ion concentration of the base of the electrolyte or the hydroxide of the base in an aqueous solution, the pH is expressed as a note. The pH is defined in the formula: $\text{pH} = -\log [\text{hydrogen ion concentration}]$. Therefore, if the pH is 13 or less, it means that the hydrogen ion concentration is 1.0×10^{-13} mol/dm³ or more. Besides, an aluminum foil having a dielectric oxide film formed on the surface at voltage of 500 V was immersed in the electrolyte (temperature 30°C). In this state, the spark ignition voltage of electrolyte (that is, the dielectric strength of electrolyte) observed by constant voltage-current elevation at constant current of 2.0 mA/cm² and the conductivity (measuring temperature 30°C) are shown.

Electrolyte A

γ -butyrolactone (100 parts), mono-1,2,3,4-tetramethyl imidazolinium phthalate (30 parts) [note: pH = 11.2], o-nitrobenzoic acid (1 part), monobutyl ester phosphate (1 part), boric acid (2 parts), and mannite (2 parts) were mixed and dissolved.

The spark voltage was 85 V, and the conductivity was 9.0 mS/cm.

Electrolyte B

γ -butyrolactone (50 parts), ethylene glycol (50 parts), trimethyl ammonium maleate (5 parts) [note: pH = 9.5], trimethylamine phthalate (5 parts) [note: pH = 9.5], diammonium adipate (3 parts) [note: pH = 9.1], boric acid (0.5 part), p-nitrobenzoic acid (1 part), and phosphoric acid (0.5 part) were mixed and dissolved.

The spark voltage was 180 V, and the conductivity was 3.3 mS/cm.

Electrolyte C

Ethylene glycol (70 parts), glycerin (30 parts), diammonium adipate (15 parts) [note: pH = 9.1], 1,6-decane dicarboxylic acid (1 part) [note: pH = 9.1], 1,7-octane dicarboxylic acid (1 part) (note: pH = 9.1), o-nitrophenol (1 part), and ammonium hypophosphite (1 part) [note: pH = 9.1] were mixed and dissolved.

The spark voltage was 340 V, and the conductivity was 0.9 mS/cm.

Electrolyte D:

γ -butyrolactone (100 parts) and tetramethyl ammonium phthalate (40 parts) [note: pH = 13.2] were mixed and dissolved.

The spark voltage was 79 V, and the conductivity was 11.5 mS/cm.

The sealing members of rubber used in the embodiments of the invention and comparative examples are as follows.

Sealing member A [vulcanization by peroxide]

It was vulcanized and formed by mixing 30 parts of rubber polymer composed of a copolymer of isobutylene, isoprene and divinyl benzene, 20 parts of carbon, 50 parts of inorganic filler, and 2 parts of dicumyl peroxide as a vulcanizing agent. The hardness of the sealing member after forming was measured on the surface portion at the side contacting with the capacitor element between two rubber holes for penetrating a lead, and the surface of the portion contacting with the lead wire side of the rubber hole inside. As a result, the IRHD (international rubber hardness degree) was respectively 67 IRHD and 66 IRHD.

Sealing member B [vulcanization by resin]

It was vulcanized and formed by mixing 30 parts of rubber polymer composed of a copolymer of isobutylene and isoprene, 20 parts of carbon, 50 parts of inorganic filler, and 2 parts of alkyl phenol formalin resin as vulcanizing agent. The hardness of the sealing member after forming

was measured on the surface portion at the side contacting with the capacitor element between two rubber holes for penetrating a lead, and the surface of the portion contacting with the lead wire side of the rubber hole inside. As a result, the IRHD (international rubber hardness degree) was respectively 77 IRHD and 76 IRHD.

(Embodiment 1)

An anode aluminum foil and a cathode aluminum foil were wound around an electrolytic paper containing Manila hemp fibers (density 0.55 g/cm^3 , thickness $50 \text{ }\mu\text{m}$). Thus prepared winding type aluminum electrolytic capacitor was kept at temperature of 300°C for 30 minutes, and the electrolytic paper was carbonized. Then, this capacitor element was immersed in a water-ethanol solution containing ethylene dioxythiophen and ferric sulfate, and lifted, and polymerized (10 minutes at 105°C), and this process was repeated 10 times. The solid organic conductive material layer composed of polyethylene dioxythiophen thus polymerized chemically was formed on the electrode foils and between electrode foils. Consequently, the capacitor element was washed in water, and dried. Further, electrolyte A was impregnated in this capacitor element. As a result, an aluminum electrolytic capacitor element with rated voltage of 50 V and electrostatic capacity of

390 μF was obtained. This capacitor element was put in an aluminum metallic case together with sealing member A, and the opening was sealed by curling process. Thus, an aluminum electrolytic capacitor was composed (size: $\phi 13 \text{ mm} \times \text{L } 20 \text{ mm}$).

(Embodiment 2)

A glass fiber nonwoven cloth (density 0.13 g/cm^3 , thickness $50 \mu\text{m}$), an anode aluminum foil, and a cathode aluminum foil were immersed in a water-ethanol solution containing pyrrole and ammonium persulfate, and lifted, dried and polymerized (10 minutes at 105°C), and this process was repeated three times. Thus, a chemically polymerized polypyrrole was formed. Then, washing in water and drying, the separator made conductive by the chemically polymerized polypyrrole, and the anode aluminum foil and cathode aluminum foil having the chemically polymerized polypyrrole formed on the surface were obtained. Then, through this conductive separator, the anode aluminum foil and cathode aluminum foil having the chemically polymerized polypyrrole formed on the surface were wound, and a capacitor element was formed. It was further immersed in soluble sulfonated polyaniline solution at concentration of 10 wt.%, impregnated at reduced pressure, lifted, and dried. Thus, the residual dried sulfonated polyaniline was formed

between the anode aluminum foil and cathode aluminum foil having chemically polymerized polypyrrole. Thus, the electric bonding between the electrode foils was reinforced. Still more, this capacitor element was impregnated in electrolyte A. As a result, an aluminum electrolytic capacitor element with rated voltage of 50 V and electrostatic capacity of 390 μF was obtained. This capacitor element was put in an aluminum metallic case together with sealing member A, and the opening was sealed by curling process. Thus, an aluminum electrolytic capacitor was composed (size: $\phi 13 \text{ mm} \times \text{L } 20 \text{ mm}$).

(Embodiment 3)

On a glass fiber nonwoven cloth (density 0.13 g/cm^3 , thickness $50 \mu\text{m}$), an anode aluminum foil, and a cathode aluminum foil, 7,7,8,8-tetracyanoquinodimethane complex in molten state was applied individually. Then, by cooling, the separator made conductive by the 7,7,8,8-tetracyanoquinodimethane complex, and the anode aluminum foil and cathode aluminum foil having the 7,7,8,8-tetracyanoquinodimethane complex formed on the surface were obtained. Then, through this conductive separator, the anode aluminum foil and cathode aluminum foil having the 7,7,8,8-tetracyanoquinodimethane complex formed on the surface were wound. Thus a capacitor element was formed.

It was further immersed in soluble sulfonated polyaniline solution at concentration of 10 wt.%, impregnated at reduced pressure, lifted, and dried at a temperature below the melting point of the 7,7,8,8-tetracyanoquinodimethane complex. Thus, the residual dried sulfonated polyaniline was formed between the anode aluminum foil and cathode aluminum foil having the 7,7,8,8-tetracyanoquinodimethane complex. Thus, the electric bonding between the electrode foils was reinforced. Still more, this capacitor element was impregnated in electrolyte A. As a result, an aluminum electrolytic capacitor element with rated voltage of 50 V and electrostatic capacity of 390 μ F was obtained. This capacitor element was put in an aluminum metallic case together with sealing member A, and the opening was sealed by curling process. Thus, an aluminum electrolytic capacitor was composed (size: $\phi 13$ mm \times L 20 mm).

(Embodiment 4)

An electrolytic capacitor was prepared in the same manner as in embodiment 1 except that electrolyte B was used as the electrolyte in embodiment 1 of the invention.

(Embodiment 5)

An electrolytic capacitor was prepared in the same manner as in embodiment 1 except that electrolyte C was

used as the electrolyte in embodiment 1 of the invention.

(Embodiment 6)

An electrolytic capacitor was prepared in the same manner as in embodiment 1 except that electrolyte D was used as the electrolyte in embodiment 1 of the invention.

(Embodiment 7)

An electrolytic capacitor was prepared in the same manner as in embodiment 1 except that sealing member B was used as the sealing member in embodiment 1 of the invention.

(Embodiment 8)

An electrolytic capacitor was prepared in the same manner as in embodiment 1 except that pyrrole was used instead of ethylene dioxythiophen in embodiment 1 of the invention.

(Embodiment 9)

An electrolytic capacitor was prepared in the same manner as in embodiment 1 except that aniline was used instead of ethylene dioxythiophen in embodiment 1 of the invention.

(Embodiment 10)

An electrolytic capacitor was prepared in the same manner as in embodiment 1 except that a mixture of iron p-toluene sulfonate and iron dodecyl benzene sulfonate was used instead of ferric sulfate, and that water-methanol solution was used instead of water-ethanol solution in embodiment 1 of the invention.

(Embodiment 11)

An electrolytic capacitor was prepared in the same manner as in embodiment 1 except that a mixture of ammonium persulfate and hydrogen persulfate water was used instead of ferric sulfate in embodiment 1 of the invention.

(Comparative example 1)

An anode aluminum foil and a cathode aluminum foil were wound around an electrolytic paper containing Manila hemp fibers (density 0.55 g/cm^3 , thickness $50 \text{ }\mu\text{m}$). Thus a capacitor element was prepared. This capacitor element was impregnated in electrolyte A, and an aluminum electrolytic capacitor element with rated voltage of 50 V and electrostatic capacity of $390 \text{ }\mu\text{F}$ was obtained. This capacitor element was put in an aluminum metallic case together with sealing member A. Then the opening was sealed by curling process. Thus, an aluminum electrolytic capacitor was composed (size: $\phi 13 \text{ mm} \times L 20 \text{ mm}$).

(Comparative example 2)

An electrolytic capacitor was prepared in the same manner as in embodiment 1 except that electrolyte A was not impregnated in embodiment 1 of the invention.

(Comparative example 3)

An anode aluminum foil and a cathode aluminum foil were wound around an electrolytic paper containing Manila hemp fibers (density 0.55 g/cm³, thickness 50 μm). Thus a capacitor element was prepared. This capacitor element was kept at temperature of 300 °C for 30 minutes, and the electrolytic paper was carbonized. Then this element was immersed in 7,7,8,8-tetracyanoquinodimethane complex in molten state and impregnated at reduced pressure. By cooling, a 7,7,8,8-tetracyanoquinodimethane complex layer was formed directly between the electrodes. Thus, an aluminum electrolytic capacitor element with rated voltage of 50 V and electrostatic capacity of 390 μF was obtained. This capacitor element was put in an aluminum metallic case together with sealing member A, and the opening was sealed by curling process. Thus, an aluminum electrolytic capacitor was composed (size: φ13 mm×L 20 mm).

(Comparative example 4)

An anode aluminum foil and a cathode aluminum foil were wound around a glass fiber nonwoven cloth (density 0.13 g/cm³, thickness 50 μm), and the obtained aluminum electrolytic capacitor element was immersed in an aqueous solution of manganese nitrate, lifted, and pyrolyzed (10 minutes at 300°C), and this process was repeated 10 times, and a manganese dioxide layer which is a solid inorganic conductive material was directly formed between the electrodes. This capacitor element was impregnated in electrolyte A. Thus, an aluminum electrolytic capacitor element with rated voltage of 50 V and electrostatic capacity of 390 μF was obtained. This capacitor element was put in an aluminum metallic case together with sealing member A, and the opening was sealed by curling process. Thus, an aluminum electrolytic capacitor was composed (size: φ13 mm×L 20 mm).

(Comparative example 5)

An electrolytic capacitor was prepared in the same manner as in comparative example 1 except that carbon fabric weaving carbon fibers was used instead of the electrolytic paper in comparative example 1.

(Comparative example 6)

An electrolytic capacitor was prepared in the same

manner as in comparative example 1 except that glass fiber nonwoven cloth coated with water dispersion type colloidal graphite was used instead of the electrolytic paper in comparative example 1.

Table 1 shows results of comparison of initial characteristics (electrostatic capacity, impedance leak current) and number of shorting troubles during aging process in aluminum electrolytic capacitors in embodiments 1 to 11 of the invention and comparative examples 1 to 6.

In each material, the number of samples was 20, and the initial characteristic (except for shorting troubles) is expressed by the average of 20 samples.

Table 1

	Electro-static capacity (μF) $f=120\text{ Hz}$	Leak current (μA) at rated voltage, 2 min	Number of shorting troubles during aging	Impedance ($\text{m}\Omega$) $f=400\text{ kHz}$
Embodiment 1	390	92	0	10
Embodiment 2	390	90	0	11
Embodiment 3	370	93	0	11
Embodiment 4	393	90	0	12
Embodiment 5	385	88	0	22
Embodiment 6	380	180	0	10
Embodiment 7	390	89	0	11
Embodiment 8	390	89	0	12
Embodiment 9	384	90	0	13
Embodiment 10	360	91	0	13
Embodiment 11	380	91	0	11
Comparative example 1	395	70	0	41
Comparative example 2	360	1000 or more	15	15
Comparative example 3	385	1000 or more	18	13
Comparative example 4	385	195	2	18
Comparative example 5	380	170	0	30
Comparative example 6	390	1000 or more	17	25

As clear from Table 1, the electrolytic capacitors in embodiments 1 to 11 of the invention is extremely small in impedance as compared with the electrolytic capacitor composed only of electrolyte in comparative example 1.

In the aluminum electrolytic capacitor lowered in resistance by using carbon fabric weaving carbon fibers as means of making the separator conductive in comparative example 5, the impedance is improved as compared with comparative example 1, but the impedance is larger than in the electrolytic capacitors in embodiments 1 to 11.

In the electrolytic capacitors in the comparative examples, that is, in the aluminum electrolytic capacitor not having electrolyte using only conductive high polymer (polyethylene dioxythiophen layer) in comparative example 2, in the aluminum electrolytic capacitor not having electrolyte using only organic semiconductor (7,7,8,8-tetracyanoquinodimethane complex layer) in comparative example 3, and in the aluminum electrolytic capacitor lowered in resistance by applying water-dispersion type colloidal graphite as the conductive means of separator in comparative example 6, shorting troubles (shorting between electrodes due to lack of dielectric strength) occurred in all samples during aging process for applying a direct voltage of 63 V at temperature of 85°C.

In the aluminum electrolytic capacitor using manganese

dioxide which is a solid inorganic conductive material, instead of solid organic conductive material, in comparative example 4, although the impedance is excellent, since the conductive material is inorganic, the electrolyte, which is an organic matter, is hardly diffused, and the restoration performance of the dielectric oxide film was not assured sufficiently. Accordingly, slight shorting troubles occurred during aging.

As explained herein, the invention presents, in a simple process, electrolytic capacitors having excellent characteristics, such as excellent impedance characteristic, small leak current, superior reliability, and high dielectric strength. Of the embodiments of the invention, in embodiment 6 of the invention using electrolyte D, since the spark ignition voltage of electrolyte is below 80 V, the dielectric strength is not sufficient, and the leak current value tends to be higher as compared with other embodiments although shorting did not occur during aging. Therefore, to express the effects of the invention sufficiently in the aspects of dielectric strength and leak current, the spark ignition voltage of the electrolyte is preferred to be 80 V or more.

Of the embodiments of the invention, in embodiment 5 of the invention using electrolyte C, since the conductivity

of the electrolyte is less than 1.0 mS/cm, the conductivity is not sufficient, and the impedance tends to be higher as compared with other embodiments. In order to express the effects of the invention sufficiently in the aspect of impedance performance, the conductivity of the electrolyte is preferred to be 1.0 mS/cm or more.

Table 2 shows the results of observation of the sealing member surface after 1000 hours of continuous application test of rated voltage of 50 V in the atmosphere of temperature of 60 °C and relative humidity of 95%, in aluminum electrolytic capacitors in embodiments 1 to 11 of the invention. The number of samples is 20 each.

Table 2

	Appearance of sealing member surface after 1000 hours of humidity resistance test at 60°C, 95% RH
Embodiment 1	No abnormality
Embodiment 2	No abnormality
Embodiment 3	No abnormality
Embodiment 4	No abnormality
Embodiment 5	No abnormality
Embodiment 6	Electrolyte leak in 2 samples
Embodiment 7	No abnormality
Embodiment 8	No abnormality
Embodiment 9	No abnormality
Embodiment 10	No abnormality
Embodiment 11	No abnormality

As clear from Table 2, extreme abnormality was not observed in any embodiment. However, in embodiment 6 using electrolyte D, since the base of the electrolytic material is a base of strong basicity with hydrogen ion concentration of less than 1.0×10^{-13} mol/dm³, in the long-term test in complex environments of high temperature and high humidity, the sealing member (sealing portion) is likely to be damaged by the effects of the base of strong basicity, and, as a result, electrolyte leak was observed

in two samples. The electrolyte leaks from the lead. In order to express the effects of the invention sufficiently in the aspect of reliability, in the base for composing the electrolytic substance, the concentration of the base or the hydroxide of the base is preferred to be 1 wt.% or more, and when the measuring temperature is 30°C, the hydrogen ion concentration of the base or the hydroxide of the base in aqueous solution is preferred to be selected at 1.0×10^{-13} mol/dm³ or more.

According to the method of embodiments 1 to 11 of the invention and comparative examples 2 and 3, again, aluminum electrolytic capacitor elements with rated voltage of 6.3 V and electrostatic capacity of 1000 μF were obtained. This capacitor element was put in a aluminum metallic case together with the sealing member A. Then the opening was sealed by curling process. Consequently, a resin seat plate made of polyphenylene sulfide was mounted. As a result, an aluminum electrolytic capacitor of vertical surface mounting type was fabricated (size: φ10 mm × L 10 mm). Thus composed aluminum electrolytic capacitor of surface mounting type was mounted on a glass epoxy substrate (2 mm thick) by using cream solder (Sn-Pb eutectic composition). By passing through a reflow furnace for infrared and hot air treatment (peak temperature 240°C, exposure time to temperature of 200 °C or more of 50

seconds), mounting and heat resistance test was conducted. The number of samples was 20.

As a result, the surface mounting type aluminum electrolytic capacitors composed in the methods conforming to embodiments 1 to 11 of the invention were suppressed in pressure elevation in the capacitor due to water adsorbed on the members because the electrolyte was composed of an organic solvent with boiling point of 200°C or more added to solid organic conductive material as electrolytic substance. Hence, mounting troubles due to scatter of sealing member of swelling of sealing member did not occur. On the other hand, in the surface mounting type aluminum electrolytic capacitors composed in the methods conforming to comparative examples 2 and 3, the pressure elevation in the capacitor due to adsorbed water was extreme, and the sealing members scattered in all 20 samples. Evidently, the surface mounting type aluminum electrolytic capacitors manufactured in the methods of the embodiments of the invention were enhanced in heat resistance when mounting, by using the electrolyte composed of organic solvent of boiling point of 200°C or more, in addition to the solid organic conductive material.

Thus, the electrolytic capacitors of the invention have excellent characteristics such as excellent impedance

WHAT IS CLAIMED IS:

1. An electrolytic capacitor comprising:

(a) a capacitor element having a positive electrode, a negative electrode, and a solid organic conductive material disposed between said positive electrode and said negative electrode,

(b) an electrolyte,

(c) a case for accommodating said capacitor element and said electrolyte, and

(d) a sealing member disposed to cover the opening of said case.

2. An electrolytic capacitor of claim 1, wherein said positive electrode comprises a metal foil and a dielectric oxide film formed on the surface of said metal foil, and has a valve action.

3. An electrolytic capacitor of claim 1, wherein said case is formed of a tubular metal having a bottom.

4. An electrolytic capacitor of claim 1, wherein said solid organic conductive material has an organic semiconductor.

5. An electrolytic capacitor of claim 1, wherein said

solid organic conductive material has a conductive polymer.

6. An electrolytic capacitor of claim 1, wherein said solid organic conductive material has an organic semiconductor and a conductive polymer.

7. An electrolytic capacitor of claim 4, wherein said organic semiconductor has at least one of 7,7,8,8-tetracyanoquinodimethane complex and its derivatives.

8. An electrolytic capacitor of claim 5, wherein said conductive polymer has a polymer of at least one monomer selected from the group consisting of pyrrole, aniline, thiophen, ethylene dioxythiophen, sulfonated aniline, sulfonated pyrrole, sulfonated thiophen, sulfonated ethylene dioxythiophen, and their derivatives.

9. An electrolytic capacitor of claim 5, wherein said conductive polymer has at least one polymer selected from the group consisting of liquid-phase chemical polymerization composition, vapor-phase polymerization composition, liquid-phase electrolytic polymerization composition, and residual dry polymer of soluble polymer solution.

10. An electrolytic capacitor of claim 5, wherein said conductive polymer has at least one polymer formed by chemically polymerizing selected from the group consisting of polypyrrole, polyethylene dioxythiophen, and polyaniline.

11. An electrolytic capacitor of claim 5, wherein said conductive high polymer has at least one polymer formed by electrolytically polymerizing selected from the group consisting of polypyrrole, polyethylene dioxythiophen, and polyaniline.

12. An electrolytic capacitor of claim 5, wherein said conductive polymer has a residual dry polymer of soluble sulfonated polyaniline solution.

13. An electrolytic capacitor of claim 1, wherein said solid organic conductive material has at least one of organic semiconductor and conductive polymer, and a residual dry polymer of soluble sulfonated polyaniline solution.

14. An electrolytic capacitor of claim 1, wherein said electrolyte has organic solvent, and at least one electrolytic substance of organic salt and inorganic salt dissolved in said organic solvent.

15. An electrolytic capacitor of claim 14, wherein said solid organic conductive material is swollen in said organic solvent.

16. An electrolytic capacitor of claim 14, wherein said electrolyte has an electrolytic substance, and the base component of said electrolytic substance has a hydrogen ion concentration of 1.0×10^{-13} mol/dm³ or more in 1 wt.% aqueous solution of the base or hydroxide of the base.

17. An electrolytic capacitor of claim 14, wherein said electrolyte has at least one selected from the group consisting of a compound having alkyl subsistent amidine group, quaternary salt of compound having alkyl substituent amidine group, tertiary amine, and ammonium.

18. An electrolytic capacitor of claim 17, wherein said electrolyte has quaternary salt of compound having alkyl subsistent amidine group, the alkyl substituent of said quaternary salt of the compound having alkyl subsistent amidine group is one of alkyl group and aryl alkyl group with 1 to 11 carbon atoms, and the compound having the amidine group of said quaternary salt of the

compound having alkyl substituent amidine group is at least one selected from the group consisting of imidazole compound, benzoimidazole compound, and alicyclic amidine compound.

19. An electrolytic capacitor of claim 14, wherein said electrolyte has at least a quaternary salt of the compound having alkyl substituent amidine group selected from the group consisting of:

1-methyl-1,8-diazabicyclo [5,4,0] undecene-7,
1-methyl-1,5-diazabicyclo [4,3,0] nonene-5,
1,2,3-trimethyl imidazolinium,
1,2,3,4-tetramethyl imidazolinium,
1,2-dimethyl-3-ethyl-imidazolinium,
1,3,4-trimethyl-2-ethyl imidazolinium,
1,3-dimethyl-2-heptyl imidazolinium,
1,3-dimethyl-2-(-3' heptyl) imidazolinium,
1,3-dimethyl-2-dodecyl imidazolinium,
1,2,3-trimethyl-1,4,5,6-tetrahydropyrimidium,
1,3-dimethyl imidazolium,
1-methyl-3-ethyl-imidazolium, and
1,3-dimethyl benzoimidazolium.

20. An electrolytic capacitor of claim 14, wherein said electrolyte has at least one property selected from

the group consisting of:

- (1) boiling point is 200°C or more,
- (2) electric conductivity at measuring temperature of 30°C is 1.0 mS/cm or more, and
- (3) spark ignition voltage is 80 V or more.

21. A manufacturing method of electrolytic capacitor comprising the steps of:

- (a) fabricating a positive electrode,
- (b) fabricating a negative electrode,
- (c) forming a solid organic conductive material on the surface of said positive electrode, and
- (d) disposing an electrode between said positive electrode having said solid organic conductive material and said negative electrode.

22. A manufacturing method of electrolytic capacitor of claim 21, wherein at said step (c), said solid organic conductive material is at least one of organic semiconductor and conductive polymer.

23. A manufacturing method of electrolytic capacitor of claim 21, wherein at said step (c), a solution containing a polymerizable monomer is bonded to the surface of said positive electrode, and said bonded monomer is

polymerized to form said solid organic conductive material.

24. A manufacturing method of electrolytic capacitor of claim 21, wherein said solid organic conductive material has at least one organic semiconductor of 7,7,8,8-tetracyanoquinodimethane complex and its derivatives.

25. A manufacturing method of electrolytic capacitor of claim 21, wherein at said step (c), a solution containing at least one monomer selected from the group consisting of pyrrole, aniline, thiophen, ethylene dioxythiophen, sulfonated aniline, sulfonated pyrrole, sulfonated thiophen, sulfonated ethylene dioxythiophen, and their derivatives is applied on the surface of said positive electrode, and said applied monomer is polymerized to form said solid organic conductive material.

26. A manufacturing method of electrolytic capacitor of claim 21, wherein at said step (c), a solution containing a polymerizable monomer is applied on the surface of said positive electrode, and said applied monomer is chemically polymerized in liquid phase to form said solid organic conductive material.

27. A manufacturing method of electrolytic capacitor

of claim 21, wherein at said step (c), said a polymerizable monomer is brought into contact with the surface of said positive electrode in a vapor-phase atmosphere of said polymerizable monomer, and polymerized in vapor phase to form said solid organic conductive material.

28. A manufacturing method of electrolytic capacitor of claim 21, wherein at said step (c), said positive electrode is immersed in a liquid having a polymerizable monomer, said monomer is electrolytically polymerized to form said solid organic conductive material on the surface of said positive electrode.

29. A manufacturing method of electrolytic capacitor of claim 21, wherein at said step (c), said solid organic conductive material of at least one of organic semiconductor and conductive polymer is formed, then said positive electrode forming said solid organic conductive material is immersed in a soluble polymer solution, and then it is dried and a residual dry polymer of said soluble polymer solution is formed on the surface of said solid organic conductive material.

30. A manufacturing method of electrolytic capacitor

of claim 21, wherein said solid organic conductive material is in a state swollen in said electrolyte.

31. A manufacturing method of electrolytic capacitor of claim 21, wherein said organic conductive material has at least one organic semiconductor of 7,7,8,8-tetracyanoquinodimethane complex and its derivatives.

32. A manufacturing method of electrolytic capacitor of claim 21, wherein said solid organic conductive material has a polymer formed from at least one monomer selected from the group consisting of pyrrole, aniline, thiophen, ethylene dioxythiophen, sulfonated aniline, sulfonated pyrrole, sulfonated thiophen, sulfonated ethylene dioxythiophen, and their derivatives.

ABSTRACT OF THE DISCLOSURE

A electrolytic capacitor includes

(a) a capacitor element having a positive electrode, a negative electrode, and a solid organic conductive material disposed between the positive electrode and the negative electrode,

(b) an electrolyte,

(c) a case for accommodating the capacitor element and the electrolyte, and

(d) a sealing member disposed to cover the opening of the case.

The solid organic conductive material has at least one of organic semiconductor and conductive polymer. In this constitution, an electrolytic capacitor having excellent impedance characteristic, small current leak, excellent reliability, and high dielectric strength is obtained.

Selected drawing: Fig. 1

Fig. 1 (a)

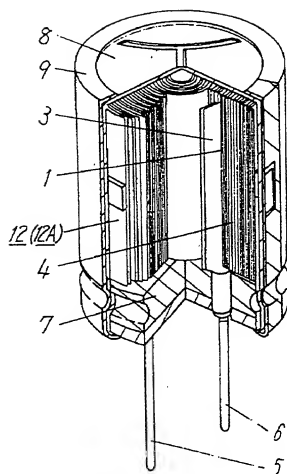


Fig. 1 (b)

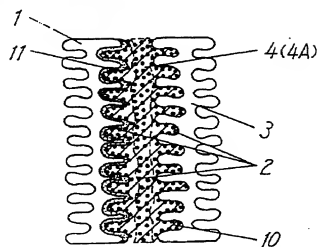
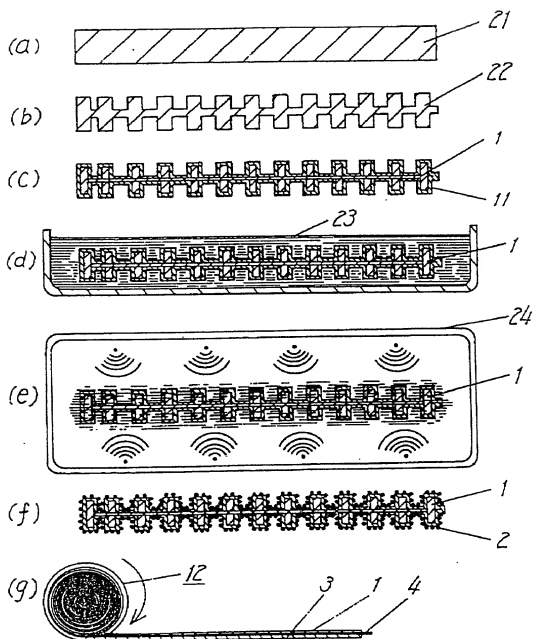


Fig. 2



P17460

Attorney Docket No. 43890-347**COMBINED DECLARATION/POWER OF ATTORNEY FOR PATENT APPLICATION**

As a below named inventor(s), I(we) hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled ELECTROLYTIC CAPACITOR AND ITS MANUFACTURING METHOD, the specification of which

(check one) ☐ is attached hereto.

☒ was filed on January 20, 1998 as
United States Application No. 09/233,936

☐ PCT International Patent Application Number _____
filed _____
and was amended on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, § 1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, § 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

Prior Foreign Application(s)

Priority Claimed

10-015269 Japan 28 January 1998
(Number) (Country) (Day/Month/Year Filed)

☒ Yes ☐ No

10-350072 Japan 09 December 1998
(Number) (Country) (Day/Month/Year Filed)

☒ Yes ☐ No

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, § 1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

(Appn. Serial No.)

(Filing Date)

(Status-patented, pending, abandoned)

(Appn. Serial No.)

(Filing Date)

(Status-patented, pending, abandoned)

I hereby appoint as my attorneys, with full power of substitution and revocation, to prosecute the patent application identified above and to transact all business in the U.S. Patent and Trademark Office connected therewith: Raphael V. Lupo (Reg. No. 28,363); Jack Q. Lever, Jr. (Reg. No. 28,149); Kenneth L. Cage (Reg. No. 26,151); Michael E. Fogarty (Reg. No. 36,139); Brian E. Ferguson (Reg. No. 36,801); Robert W. Zelnick (Reg. No. 36,976); Edward E. Kubasiewicz (Reg. No. 30,020); Paul Devinsky (Reg. No. 28,553); and Wilhem F. Gadiano (Reg. No. 37,136); Laura A. Donnelly (Reg. No. 38,435); Craig L. Plastrik (Reg. No. 41,254); David A. Spenard (Reg. No. 37,449)

Please address all correspondence and telephone calls to:

Edward E. Kubasiewicz
MCDERMOTT, WILL & EMERY
600 13th Street, N.W.
WASHINGTON, D.C. 20005-3086
Telephone: 202-756-8000

**020277**

PATENT AND TRADEMARK OFFICE

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of sole or first inventor **Kazuyo SAITO**

Inventor's signature Kazuyo Saito Date April 5, 1999
Residence **6-1-215, Uyamahigashimachi, Hirakata-shi, Osaka 573-1136 JAPAN**
Post Office Address SAME AS ABOVE Citizenship Japanese

Full name of second inventor **Yukihiko NITTA**

Inventor's signature Yukihiko Nitta Date April 5, 1999
Residence **2-1-311, Nishifura, Kohata, Uji-shi, Kyoto, 611-0002, JAPAN**
Post Office Address SAME AS ABOVE Citizenship Japanese

Full name of third inventor **Hiroshi TADA**

Inventor's signature Hiroshi Tada Date April 5, 1999
Residence **3-40-22, Noda, Kuzuha, Hirakata-shi, Osaka, 573-1103 JAPAN**
Post Office Address SAME AS ABOVE Citizenship Japanese

Full name of fourth inventor **Shigeyoshi IWAMOTO**

Inventor's signature Shigeyoshi Iwamoto Date April 5, 1999
Residence **95-36, Hijiri, Tefada, Joyo-shi, Kyoto, 610-0121 JAPAN**
Post Office Address SAME AS ABOVE Citizenship Japanese